CLAIMS

- Method of assaying nucleic acids by molecular hybridization , characterized by the fact
 that it includes the following step :
 - taking samples of biological material by a sampling device (1; 13) comprising abrasive sampling means (2;15, 16) capable of retaining biological material in the form of cells.
- Method according to Claim 1, characterized by the fact that the sampling of biological
 material is done in the surrounding air.
 - 3. Method according to any of claims 1 or 2, characterized by the fact that the sampling is done outside of a laboratory where the assaying will be done, and in that the method includes a step of transport of the abrasive sampling means (2;15,16) loaded with their respective samples of biological material to said laboratory.
- 15 4. Method according to any of the preceding claims, characterized by the fact that it furthermore includes a step of extraction of the nucleic acids, comprising
 - an immersion step into an extraction buffer of the abrasive sampling means (2;15,
 16) loaded with their respective samples of biological material
 - a step of agitation in the extraction buffer
- 20 a separation step, and

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- a recovery step for recovering clarified solution containing the nucleic acids.
- 5. Method according to Claim 4, characterized by the fact that the separation step consists in a centrifugation, and the supernatant constitutes the clarified solution.
- 6. Method according to any of the preceding claims, characterized by the fact that the assaying by molecular hybridization is done by polymerase chain reaction (PCR).
- 7. Method according to any of the preceding claims, characterized by the fact that the assaying of nucleic acids by molecular hybridization is done in order to determine the presence of a pathogenic agent in the biological material.
- 8. Method according to any of the preceding claims, characterized by the fact that the biological material consists of material of plant origin.
- 9. Kit for implementing the method according to any of claims 1 to 8, characterized by the fact that it comprises a sampling device (1; 12) comprising abrasive sampling means (2; 14,15) able to retain biological material in the form of cells.
- 10. Kit according to Claim 9, characterized by the fact that the sampling means (2; 16, 17) comprise a solid material having an abrasive outer surface.

- 11. Kit according to Claim 11, characterized by the fact that the solid material is chosen from the group consisting of silica, glass, metals, carbon fibers and plastics.
- 12. Kit according to any of claims 10 and 11, characterized by the fact that the abrasive outer surface comprises asperities able to retain cells of biological material.
- 5 13. Kit according to any of claims 9 to 12, characterized by the fact that the sampling device (2, 13) comprises a support (3,5,6;14,17) able to support the abrasive sampling means (2; 15,16).
 - 14. Kit according to any of claims 9 to 13, characterized by the fact that it comprises means (11) for the transport of the abrasive sampling means (2; 15, 16)
- 15. Kit according to any of claims 9 to 14, characterized by the fact that it comprises means of identification of the abrasive sampling means.
 - 16. Kit according to any of claims 9 to 15, characterized by the fact that it comprises extraction buffer for assaying nucleic acids by hybridization.
- 17. Kit according any of the preceding claims, characterized by the fact that it comprises specific reagents of PCR reactions.